Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

Claim 1 (canceled).

Claim 2 (currently amended): The method according to claim ± 27 , wherein the donor cells (6) contain naturally occurring stem cells.

Claim 3 (currently amended): The method according to claim ± 27 , wherein the cells (2) of the morula (7) or the internal cell mass (4) of the blastocyst (1) are prepared in a culture dish (8, 9, 10) or are used to prepare a soluble matrix fraction.

Claim 4 (currently amended): The method according to claim ± 27 , wherein the donor cells (6) are obtained from umbilical cord blood.

Claim 5 (currently amended): The method according to claim ± 27 , wherein the donor cells (6) are obtained from placenta.

Claim 6 (currently amended): The method according to claim $\frac{1}{27}$, wherein the donor cells $\frac{1}{6}$ are obtained from bone marrow.

Claim 7 (currently amended): The method according to claim $\frac{1}{27}$, wherein the donor cells $\frac{1}{6}$ are obtained from fatty tissue.

Claim 8 (currently amended): The method according to claim $\frac{1}{27}$, wherein the cells $\frac{(2)}{(2)}$ of the morula $\frac{(7)}{(7)}$ or the internal cell mass $\frac{(4)}{(7)}$ of the blastocyst $\frac{(1)}{(7)}$ are tetraploid cells.

Claim 9 (currently amended): The method according to claim † 27, wherein the cells (2) of the morula (7) or the internal cell mass (4) of the blastocyst (1) has cells whose genome contains vectors that cause a lethal sensitivity to appropriate cultivation conditions in comparison to the particular corresponding wild type.

Claim 10 (currently amended): The method according to claim \pm 27, wherein the genome of the donor cells (6) contains a vector which causes a resistance to additives of culture media.

Claim 11 (currently amended): The method according to claim ± 27 , wherein the survivability of the cells (2) of the morula

(7) or the internal cell mass (4) of the blastocyst (1) is reduced by adding suitable selected antibodies.

Claim 12 (currently amended): The method according to claim 9, wherein the survivability of the cells (2) of the morula (7) or the cells of the internal cell mass (4) of the blastocyst (1) is reduced in a way that is tailored to the varying degrees of differentiation of the donor cells (6) and is chronologically well-ordered.

Claim 13 (currently amended): The method according to claim \pm 27, wherein before the donor cells (6) are supplied into the morula (7) or the blastocyst (1), the donor cells (6) are brought into contact in culture dishes with other blastocysts or internal cell masses isolated from other blastocysts, and those donor cells having a relatively high contact affinity are isolated and supplied to the morula (7) and/or blastocyst (1) first cited.

Claim 14 (currently amended): The method according to claim ± 27 , wherein before the donor cells (6) are supplied into the morula (7) or the blastocyst (1), the donor cells (6) are equipped with a genetic marker that ensures cells having a lower degree of differentiation are isolated and supplied into the morula (7) or blastocyst (1).

Claim 15 (currently amended): The method according to claim ± 27 , wherein the morula (7) or blastocyst (1) is a mouse morula or mouse blastocyst.

Claim 16 (currently amended): The method according to claim ± 27 , wherein the morula (7) or blastocyst (1) is a pig morula or pig blastocyst.

Claim 17 (currently amended): The method according to claim ± 27 , wherein when the donor cells (6) are supplied to a blastocyst (1), the supply is performed through injection.

Claim 18 (currently amended): The method according to claim ± 27 , wherein when the donor cells (6) are supplied to a morula (7), the supply is performed through aggregation.

Claim 19 (currently amended): The method according to claim ± 27 , wherein the donor cells (6) are human donor cells.

Claims 20-22 (canceled).

Claim 23 (currently amended): The method according to claim \pm 27, wherein the donor cells (6) are donor cells of non-human mammals.

Claims 24-26 (canceled).

Claim 27 (new): A method for producing cell lines or individual organs comprising the steps of:

- (a) cultivating a nonhuman morula or a nonhuman blastocyst under conditions that enable a further development of the morula or blastocyst to occur in stages in which newly formed cell lines having a high degree of differentiation are produced;
- (b) supplying differentiable donor cells to the morula or the blastocyst to produce cell lines; and
- (c) isolating the cell lines or further differentiating the cell lines into organs through transfer of the blastocyst into a surrogate mother animal;

wherein the cells of the morula or an internal cell mass of the blastocyst have a restricted survivability in comparison to a corresponding wild type or survivability of the cells or the internal cell mass is reduced through selected cultivation conditions; and

wherein the donor cells supplied to the morula or blastocyst have varying degrees of differentiation and are of non-embryonic origin.

ELECTION OF INVENTION:

The Patent Examiner has made new grounds of restriction and requires the selection of one of the following groups of invention for further prosecution:

Group I: Claims 1-19 and 23 (claims 1-4, 8-14, 16, 17 and 19, drawn to elected species), drawn to a method for producing cell lines; or

Group II: Claims 1-19 and 23 (claims 1-4, 8-14, 16, 17 and 19, drawn to elected species), drawn to a method for producing individual organs.

ELECTION:

Applicants respectfully elect, with traverse, the invention of Group I, claims 27 (claim 1 having been canceled), 2-19 and 23, for further prosecution.